

***In-Vitro* Alpha Amylase Inhibition and Antioxidant Activities of Leaves Extract of Sundanese Traditional Salad (*Lalapan*) from Indonesia**

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ABSTRACT

This research aims to evaluate the α -amylase inhibitory activity and antioxidant potential of leaves extract of Gandaria (*Bouea macrophylla* Griff.), Basil (*Ocimum africanum* Lour.), Pohpohan (*Pilea melastomoides* (Poir.) Wedd.), and Lettuce (*Lactuca sativa* L.) as common vegetables used in Sundanese traditional salad. Extraction methods used were traditional and maceration methods. Phytochemical screening was used to determine the phytochemical component qualitatively. Quantitative estimation of phenols (Folin-Ciocalteu method), flavonoids (AlCl_3 colorimetric assay), total antioxidant activity along with free radical scavenging activity (DPPH) was also carried out. Soluble starch was used as substrate for analysis of α -amylase inhibitory activity, which is calculated by IC_{50} value. Phytochemical screening revealed the presence of phytochemical constituents like flavonoids, tannins, saponins, and polyphenols compound in both extraction methods. The results showed that the value of Total Phenolic Content (TPC), Total Flavonoid Content (TFC), antioxidant activity, and α -amylase inhibitory in both extraction were significantly different for each leaves extracts ($p < 0.05$). The gandaria leaves extracted with maceration method had the highest value among the leaves analysed. The value of TPC, TFC, antioxidant activity were 364.56 ± 65.97 mg Gallic Acid Equivalent (GAE)/g extract, 70.2 ± 10.54 mg Quercetin (QE)/g extract, and $35 \mu\text{g/ml}$ of IC_{50} , respectively. The extract of gandaria maceration leaves exhibited significant α -amylase inhibitory activity with an IC_{50} value of $60 \mu\text{g/ml}$. Various pharmacologically active compounds were found in the local vegetables, thus they can be rich sources of antioxidants and α -amylase inhibitor activity. Hence, could be developed as vegetables based functional food products.

Keywords: α -amylase, antioxidant, extraction methods, *lalapan*

INTRODUCTION

Diabetes mellitus is one of the diseases associated with an increased production of Reactive Oxygen Species (ROS) and impaired antioxidant defense system which result in oxidative damage (Kundusen *et al.* 2011). Diabetes mellitus is also characterized by hyperglycemia with disturbances of carbohydrate, lipid, and protein metabolism resulting from defect in insulin secretion and/or insulin action (Keerthana *et al.* 2013). Thus reducing the postprandial hyperglycemia to lower glucose absorption using inhibitors of carbohydrate digesting enzymes is important (Bhuvaneshwari *et al.* 2014). The α -amylase is an enzyme to break down starch into simple sugars, such as dextrin, maltotriosa, maltose, and glucose. Inhibiting the activity of α -amylase

enzyme is an effective method to control blood glucose (Alexander & Maltodextrins 1992). Bioactive components found in herbal plants have shown good antioxidant and anti-diabetic properties (Kunyanga *et al.* 2012). As a bioactive compound, phytochemicals components such as tannins, phenols, and flavonoids have potential as inhibitors of the α -amylase enzyme (Keerthana *et al.* 2013).

Blood glucose can be controlled by maintaining diet or dietary behavior, regular physical activity, maintaining body weight, and consuming vegetables that can help regulate blood glucose concentration effectively (Prabhakar & Doble 2008). *Lalapan* (Sundanese traditional salad) is a raw and fresh vegetable dish commonly consumed by Sundanese in their daily meal. Pratama (2014), identified that the most common vegetables served as part of this traditional salad

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were basil, pohpohan, and lettuce leaves. Another leave commonly consumed by Sundanese and is containing phytochemical compounds and health benefits is gandaria leaves (Lolaen *et al.* 2013; Rajan & Bhat 2016). Previous research on antioxidant activity showed that the leaves of gandaria, basil, pohpohan, and lettuce have a high antioxidant activity (Andarwulan *et al.* 2010; Marwat *et al.* 2011; Gan & Azrina 2016; Andina & Musfirah 2017).

Considering the local availability and eating habit as well as findings on the potential health benefits of these leaves, it is important to explore the bioactive components potential for health. The result can inform the development of functional food based on locally available vegetables. The exploration of the leaves bioactive components should also consider the extraction method since the extraction method will affect the extracted compound. There are two extraction methods according to Verawati *et al.* (2016), they are the traditional and the laboratory or maceration methods. The traditional and the maceration methods are often used because they are relatively easy to conduct and do not require special tools. However, studies have shown that extraction processes using either the traditional or the maceration methods affected the total phenolic compound extracted from the leaves (Verawati *et al.* 2016). Therefore, it is imperative to identify the antioxidants and α -amylase inhibitory activity of the Sundanese traditional salad (*Lalapan*) using both the traditional and the maceration extraction method.

METHODS

Design, location, and time

This experimental study used a randomized block design. The research activities included sample extraction, phytochemical screening, analysis of phenol and flavonoid, antioxidant activity, and inhibition of α -amylase carried out in the Laboratory of Chemistry and Food Analysis, Department of Community Nutrition, IPB University, started from October–January 2019.

Materials and tools

The gandaria leaves was collected from Indonesia Institut of Science, Cibinong Bogor, while basil, pohpohan, and lettuce leaves

were collected from Cigombong Agricultural Technology Park, Bogor. Specimens have been authenticated (No:2115a/IPH.1.01/If.07/XII/2018) with the following results are Gandaria (*Bouea macrophylla* Griff.), Basil (*Ocimum africanum* Lour.), Pohpohan (*Pilea melastomoides* (Poir.) Wedd.), and Lettuce (*Lactuca sativa* L.).

Materials used for extraction and analysis were water for solvent, aquades, container, Wagner reagents, Liebermann-Burchard reagent, HCl, methanol, Mg, KOH, FeCl₃, gallic acid, Folin-Ciocalteu, Na₂CO₃, quercetin, AlCl₃, NaNO₂, NaOH, vitamin C, 1.1-diphenyl-2-picrylhydrazyl (DPPH), porcine pancreatic α -amylase enzyme concentration of 30 U/mg, 1% starch, phosphate buffer pH 6.9, 3.5-dinitrosalicylic solution (DNS), and potassium sodium tartrate.

The tools used for extraction and analysis were analytical scale, knife, board, filter cloth, Whatman 41, Buchner, blender, centrifuge, hot plate, and vacuum evaporator, erlenmeyer, test tubes, micro pipet, cuvettes, vortice, water bath, and UV-VIS spectrophotometers (Shimadzu UV-160A, Kyoto, Japan).

Data collection

Sample preparation and extraction of leaves material. Sorted leaves were dried in shade at room temperature to choose the edible parts, leaves were washed in tap water and drained at room temperature, then leaves were chopped into small pieces to obtain a larger surface area. The extraction method used were traditional extraction and the maceration (laboratory). In the traditional extraction, samples were blended with water (1:10 w/v), then centrifuged at 2,000 rpm (5 minutes) and the extract was filtered through a Whatman 41 using Buchner, and evaporated using a hot plate to obtain the liquid extract (Aisyah *et al.* 2015 modification). While in the maceration method, the sample was placed in a container and soaked in water. The container with its contents was sealed and kept for 2x24 hours with some stirring. Subsequently, the entire mixture then filtered by a filter cloth, then filtered again through Whatman 41 using Buchner. After filtration process, the macerate was evaporated at 60°C (65 kPa) using vacuum evaporator to obtain the dry extract (Verawati *et al.* 2016 modification).

The evaporation process of traditional extracts aimed to make it easier to control the

concentration of the solution. Stock solution was done by weighing both extracts obtained from the two different method in the same amount. Thus, it was expected that the Total Dissolve solid (TDS) in the stock solution could be controlled despite the amount was not exactly the same due to the different sample form.

Phytochemical screening. The extracts were analyzed for the active phyto-constituents such as alkaloids, flavonoids, tannin, saponin, polyphenols, and terpenoid according to Harborne (1987). Alkaloids were analyzed by dissolving 2 ml of sample in 2 ml of 2% HCl, then heated for 5 minutes and filtered. The filtrate was added by 2–3 drops of Wagner reagent. The formation of white deposits shows the presence of alkaloid compounds. Flavonoids were analyzed by dissolving 2 ml of sample in 2 ml of methanol, then added with Mg and 5 drops of 37% HCl. The formation of red or orange colour indicates the presence of flavonoids. Tanin was analyzed by adding 2 ml of sample with FeCl_3 reagent. The formation of dark blue or blackish green colour indicates the presence of tannin. Saponin was analyzed by dissolving 2 ml of sampel in a test tube, 10 drops of KOH were added and then heated in a water bath (50°C) for 5 minutes and the solution was shaken for 15 minutes. The presence of a saponin is arranged with a steady foam and stable for 15 minutes. Polyphenols were analyzed by dissolving 2 ml of sample in 10 ml distilled water and heated for 5 minutes, then filtered. The filtrate was added with 4–5 drops of FeCl_3 5%. The formation of dark blue or blackish green colour indicates the presence of a phenols. Terpenoid was analyzed by adding 2 ml of sample with 1 ml of the Liberman-Burchard reagent. The formation of dark blue or blackish green colour indicates the presence of terpenoid compounds.

Estimation of total phenols. One ml of the extract solution was mixed with 1 ml of Folin-Ciocalteu reagent, then 1 ml of aqueous 10% Na_2CO_3 solution was added and vortexed. The mixture were allowed to stand for 90 minutes at room temperature. Absorbance was measured using a spectrophotometer at 765 nm. The TPC was subsequently calculated using gallic acid as standard. Results obtained were expressed as mg GAE/g extract (Singleton & Rossi 1965).

Estimation of total flavonoid. One ml of the extract solution dissolved in aquades (1:4), added 60 μl of 5% NaNO_2 solution (incubated for

5 minutes), then added with 60 μl of 10% AlCl_3 solution (incubated for 6 minutes). After that, added with 400 μl of 10% NaOH solution and then prepared by aquades to 2 ml. The mixture then was vortexed. The absorbance was measured using a spectrophotometer at 510 nm. The TFC was calculated using quercetin as standard, and results obtained were expressed in term of QE/g extract (Sahu & Saxena 2013).

DPPH free radical-scavenging assay.

Antioxidant activity were determined based on percentage inhibition of DPPH radical and used vitamin C as a standard. The results obtained are presented by the IC_{50} value (concentration of the sample required to inhibit 50% of radical). The lower of IC_{50} value indicates that antioxidant activity is getting higher (Molyneux 2004). 750 μl of extracts solution at different concentrations (0.25; 0.5; 1; 2; 4 mg/ml) were added to 750 μl of 0.2 mM DPPH methanolic solution, the mixture was vortexed and left to stand in the dark for 30 minutes. Absorbance was measured at 517 nm. The percentage of inhibition of DPPH radical was calculated according to the following equation:

$$\% \text{Inhibition} = \left(\frac{\text{Abs control} - \text{Abs sample}}{\text{Abs control}} \right) \times 100$$

whereas, Abs control is the absorbance of DPPH solution without leave extracts.

In-vitro α -amylase inhibitory assay. The analysis of α -amylase inhibitory was conducted using the method described by Apostolidis *et al.* (2007) with slight modification. Acarbose as the pharmacological inhibitor was included as the positive control. Thirteen μl of extracts solution at different concentrations (0.078; 0.156; 0.3125; 0.625; 1.25; 2.5; 5; 10 mg/ml) were added to 94 μl phosphate buffer (pH 6.9) containing α -amylase solution (30 U/mg). Then, 63 μl of 1% soluble starch and 207 μl phosphate buffer (pH 6.9) were added to each tube at time intervals. The mixture was incubated at 37°C for 10 minutes. The reaction was stopped with a 313 μl of DNS reagent. The test tube were then incubated at 100°C for 10 minutes and cooled to room temperature. The mixture was then diluted with 313 μl of 40% potassium-sodium tartrate. Absorbance was measured at 540 nm. The percentage of α -amylase inhibitory was calculated according to the following equation:

$$\% \text{Inhibition} = \left(\frac{\text{Abs control} - (\text{Abs sample} - \text{Abs blank})}{\text{Abs control}} \right) \times 100$$

whereas, Abs control is the absorbance of solvent without sample, Abs blank is the absorbance of solvent without enzyme.

Data analysis

Data processing used the Microsoft Excel 2013 and Statistical Program Social Science (SPSS) version 16.0. Data analysis comprised of descriptive analysis, normality test, and different test. Different test were analyzed using Two Way ANOVA and Least Significance Differences (LSD) for differences of each samples ($p < 0.05$). Results were expressed as the mean \pm standard deviation.

RESULT AND DISCUSSION

Phytochemical screening and quantification of secondary metabolite

Qualitative phytochemical screening was aimed to identify any chemical components with bioactive compounds in the plant extracts, which can be further utilized (Aharoni & Galili 2011). The qualitative analysis of the leaves extract confirmed the presence of flavonoids, tannins, saponins, and polyphenol compounds in both of the traditional and the maceration methods (Table 1).

The results of phytochemical screening showed different result from previous research, such as for mango (*Mangifera indica*) that is in the same family as gandaria (*Anacardiaceae*), basil (*O. basilicum*), pohpohan (*Pilea trinervia*), and lettuce (*Lactuca sativa*) leaves. However it is important to note that the leaves species as well as the extraction method used in these research were varied (Adham 2015; Dhital 2017; Guntara *et al.* 2016; Algfri *et al.* 2019). Thus, difference

in the phytochemical components found in this study might be caused by several factors such as differences in the extraction steps, the polarity of solvent, and the species of plant. In addition, differences in the secondary metabolites component found in plants can be due to soil conditions where the plants are planted, such as soil moisture, water availability, temperature, pH, land conditions, and light intensity (Vinolina 2014).

Phenolic compounds are the most widely distributed secondary metabolites and play an important role in growth regulators, have antioxidant, structural, attractant, signaling and protective functions (Babenko *et al.* 2019). The value of Total Phenolic Content (TPC) in both extraction methods were significantly different for each leaves extracts ($p < 0.05$) (Figure 1). The results also showed that phenolic content also significantly different between each of leaves species, thus extraction methods and the leaves species affected the total phenolic content.

The results showed that gandaria leaves extract with maceration had the highest total phenolic content, followed by gandaria from traditional extraction, lettuce, basil, and pohpohan with maceration extraction. The leaves extracted with traditional method showed lower results of TPC compared to the maceration. This is probably that maceration extract was evaporated using a vacuum evaporator at 60°C (65 kPa), so the water content evaporates more than in the traditional evaporation using a hot plate. The final result of maceration is dry extract obtained by drying the liquid extract until specific concentration according to the procedure. Drying is the process for removing solvents from the material to obtain powder/dry extract, while traditional extraction

Table 1. Phytochemical screening of leaves extract

Phytochemicals	Traditional				Maceration			
	Gandaria	Basil	Pohpohan	Lettuce	Gandaria	Basil	Pohpohan	Lettuce
Alkaloids	-	-	-	-	-	-	-	-
Flavonoids	+++	++	++	++	+++	++	++	++
Tannin	+++	++	+	+	+++	++	+	+
Saponins	+	+	++	+	+	++	+++	+
Polyphenols	+++	++	+	+	+++	++	+	+
Terpenoids	-	-	-	-	-	-	-	-

+ Present; - Absent

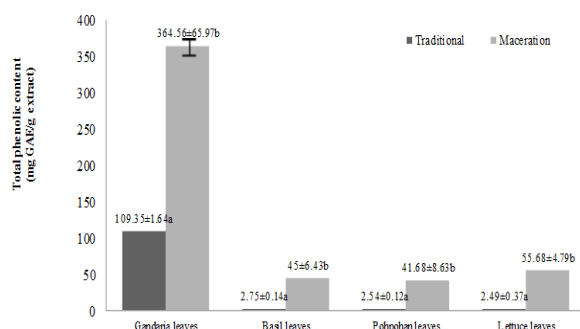


Figure 1. Total phenolic content of leaf extract. Different letters (a-b) for each leaves are significantly different ($p < 0.05$)

is liquid extract so the concentration couldn't be controlled properly (Martin *et al.* 1961).

The evaporation process uses a vacuum so that it is good for food products that are sensitive to high temperatures. It can be processed to reduce their water content without damaging the physical condition and chemical content of the material (Supriatna 2008). In traditional extraction method, the extraction results were evaporated using a hot plate at 100°C without vacuum until the volume was reduced by 50% to get the liquid extract. While in the maceration, results were evaporated using an evaporator at 60°C so that the possibility of losing the water soluble nutrients will be even lower and the average of water evaporated was 98.8% by vacuum to get dry extract.

Verawati *et al.* (2016) reported that the total phenolic content of piladang water leaves extract with traditional extraction method (*S. scutellarioides* L. Codd), evaporated by hot plate showed lower results (77.31 ± 0.21 mg GAE/g extract) compared to piladang maceration leaves extract (293.30 ± 0.13 mg GAE/g extract). Difference in extraction method affected the extracted compound. Andina and Musfirah (2017) reported that the TFC of gandaria leaves extracted by maceration was 68.53 ± 1.37 mg GAE/g extract equal to our findings with the same method. It showed that the extraction method in different process affects the secondary metabolites content. In addition to extracted methods, differences in soil conditions or growing environment of plants will also affect the content of chemical compounds in these plants. The results of secondary metabolite content will show

different on the same plant type if these plants are planted in different places (Yenni 2012).

Flavonoids are antioxidant compounds and naturally produced as secondary metabolites in plants. The Total Flavonoid Content (TFC) in different extraction methods were significantly different for each leaves extracts ($p < 0.05$) (Figure 2). The results also showed that flavonoid content also significantly different between each of the leaves extracts. It means that the extraction methods and leaves species affected the total flavonoid content of gandaria, basil, pohpohan, and lettuce leaves.

The maceration leaves extract contained higher TFC compared to the traditional leaves extract. The highest flavonoid content was found in gandaria from maceration method, followed by pohpohan, basil, and lettuce from maceration extracts. In contrast to the TPC, the gandaria traditional leaves extract showed was ranked fifth in TFC content.

Dhital's research (2017) showed that mango leaves as a plant from the same family with gandaria (*Anacardiaceae*), contained flavonoids of 55.63 mg QE/g maceration extract. It means that *Anacardiaceae* family contains high flavonoids as antioxidants. Geographical location, time and climate differences, soil conditions, administration of pesticides or herbicides in plants are factors that can influence vegetable nutrients, antioxidants, and flavonoids (Andarwulan *et al.* 2010).

Overall, the TPC and TFC in maceration leaves extract had higher values than the traditional extraction. In the maceration method the sample or material to be extracted is soaked

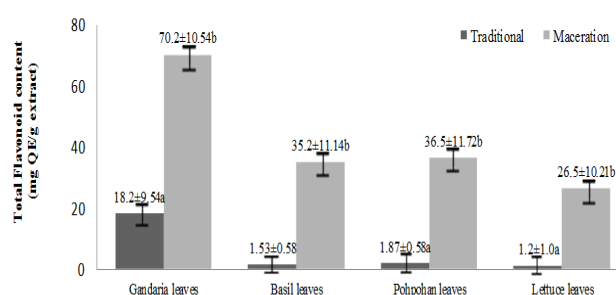


Figure 2. Total flavonoid content of leaves extract. Different letters (a-b) for each leaves are significantly different ($p < 0.05$)

in a closed container using a solvent and allowed to remain at room temperature with an adequate agitation. This process aims to soften or break down the plant cell walls so that phytochemical compounds can dissolve in solvents. During the immersion process, the cell wall and cell membrane of the sample will break because of the different pressure between the inside and outside of the cell and the content of secondary metabolites in the cell plasma will be dissolved in the solvent. In addition, the time of maceration can be adjusted according to the research purpose thus able to yield better result (Ibrahim & Marham 2013).

The selection of solvents in the extraction process will also affect the effectiveness of the traceability based on solubility of natural material compounds in the solvent (Ibrahim & Marham 2013). Solvents that have the same polarity as solutes dissolve the substance better. Water is one of the solvents that is often used in extraction processes with high polarity (Altemimi *et al.* 2017). High polarity solvents will also attract phytochemical components that also have high polarity levels such as phenols and flavonoids (Harborne 1987; Robinson 1995; Stankovic 2011).

The IC_{50} value of gandaria, basil, pohpohan, and lettuce leaves showed that all maceration leaves extracts had higher antioxidant activity compared to traditional leaves extracts, especially the gandaria leaves extract showed the highest value of antioxidant activity (Table 2). Vitamin C as a standard in this research had an IC_{50} value of 10 $\mu\text{g/ml}$. A low value of antioxidant activity could be caused by the content of bioactive components of each sample with different extraction, and crude extract without partitioning used in this research is probably cause antioxidant activity to be of low value (Rahman *et al.* 2017). Andina and Musfirah (2017) reported that the antioxidant activity of gandaria leaves extract with maceration was high in antioxidant activity (IC_{50} 55.83 $\mu\text{g/ml}$). It means that gandaria showed the highest antioxidant potential.

Our results showed that antioxidant activity in both extraction were significantly different for each leaves extracts ($p < 0.05$). The results showed that antioxidant activity also significantly different between each of leaves extracts. It means that the extraction method and leaves affected the antioxidant activity of gandaria,

basil, pohpohan, and lettuce leaves. Manggala *et al.* (2017) reported that the antioxidant activity of mango leaves extract with maceration method (*Mangifera indica*) was higher (IC_{50} 5.09 $\mu\text{g/ml}$) compared to the extract from reflux method (IC_{50} 8.35 $\mu\text{g/ml}$).

Analysis of α -amylase inhibitory using acarbose as a positive control, showed very high inhibitory activity with IC_{50} value of 0.00045 $\mu\text{g/ml}$. Our results showed that gandaria and basil leaves extract with maceration have high α -amylase inhibitory activity (IC_{50} 60 and 70 $\mu\text{g/ml}$) compared to other samples. Inhibitory action for α -amylase enzyme in both extraction were significantly different for each leaves extracts ($p < 0.05$). The results showed that α -amylase inhibitory activity also significantly different between each of the leaves species extracts. It means that the extraction methods and leaves affect the inhibitory activity of gandaria, basil, pohpohan, and lettuce leaves. Hence, we found that the leave extract which showed the highest α -amylase inhibitory activity is the one that has the highest antioxidant activity, TPC, and TFC value. The extract was gandaria leaves with maceration extraction method (Table 3).

Bhuvaneshwari *et al.* (2014) showed that mature and tender mango leaves extracted by reflux had an IC_{50} values of 22.01 and 35.73 $\mu\text{g/ml}$, respectively. It showed that the *Anacardiaceae* family has very high of α -amylase inhibitory activity. El-Beshbishy and Bahashwan (2011) reported that the inhibitory activity of α -amylase enzyme in basil extract was indicated by IC_{50} value of 42.50 mg/ml . The inhibitory activity is probably from the phenolic and flavonoid content of basil. Rahayuningsih (2014) reported that the

Table 2. Antioxidant activity of leaves extract

Leaves extract	IC_{50} ($\mu\text{g/ml}$)	
	Traditional	Maceration
Vitamin C (10 $\mu\text{g/ml}$)**		
Gandaria (<i>B. macrophylla</i>)	649 ^a	35 ^b
Basil (<i>O. africanum</i>)	>2,000 ^a	670 ^b
Pohpohan (<i>P. melastomoides</i>)	>2,000 ^a	250 ^b
Lettuce (<i>L. sativa</i>)	>2,000 ^a	183 ^b

* Different letters (a-b) for each leaves are significantly different ($p < 0.05$)

** Vitamin C as standar

Table 3. α -amylase inhibitory activity of leaves extract

Leaves extract	IC ₅₀ (μ g/ml)	
	Traditional	Maceration
Acarbose (0.00045 μ g/ml)**		
Gandaria (<i>B. macrophylla</i>)	163 ^a	60 ^b
Basil (<i>O. africanum</i>)	>1,000 ^a	70 ^b
Pohpohan (<i>P. melastomoides</i>)	>1,000 ^a	105 ^b
Lettuce (<i>L. sativa</i>)	>1,000 ^a	117 ^b

*Different letters (a-b) for each leaves are significantly different (p<0.05)

**Acarbose as positive control

flavonoid content pohpohan (*Pilea trinervia*) leaves extract had an anti-hyperglycemia effect as indicated by glucose tolerance test in glucose induced diabetic rats. Lactucaxanthin (Lxn) of lettuce (*Lactuca sativa*) function as an inhibitor of carbohydrate digestive enzymes. Gopal *et al* (2017), reported that the inhibitory activity of the α -amylase enzyme from Lxn had an IC₅₀ value of 434.5 μ g/ml.

Maintaining blood glucose levels effectively can be obtained from natural sources such as from the bioactive compounds in vegetables that can function as α -amylase inhibitor. Natural sources inhibitors, such as the one coming from the vegetables commonly consumed are highly recommended because they tend to not cause side effects when consumed (Tundis *et al.* 2010). In addition, Samudra *et al.* (2015) stated that secondary metabolites have high ability to inhibit the activity of the α -amylase enzyme compared to the primary metabolites.

Gu *et al.* (2015) stated that phenolic acids and flavonoids can inhibit the activity of the α -amylase enzyme. Phenolic acids and flavonoids will form covalent bonds with α -amylase into quinones or lactones and react with nucleophilic groups, this causes changes in the activity of the α -amylase enzyme (Tadera *et al.* 2006). Reaction of phenolic compounds and proteins be able to inhibit enzymatic activity including α -amylase enzymes (Yanishlieva-Maslarova 2001). Polyphenols will form complexes with starch so that the α -amylase enzyme cannot recognize the substrate (Himmah & Handayani 2012).

The effect of flavonoids on the α -amylase enzyme using enzymatic kinetic and fluorescence spectroscopy shows that flavonoids will form complexes with the α -amylase enzyme. Flavonoids will be bound to the α -amylase enzyme in a hydrophobic (Yuan *et al.* 2014). Flavonoids are also able to inhibit α -amylase activity because they have the ability to form quinones with 4-oxopyrane (Adisakwattana & Chanathong 2011). Flavonoids as antioxidants are able to prevent a decrease of β -pancreatic cells function progressively that caused by oxidative stress. Thus, it can reduce the occurrence of type 2 diabetes and can prevent the complications of diabetes in the long term (Song *et al.* 2005).

Within this research, the amount of Total Dissolve Solid (TDS) in samples with different extractions was not exactly the same thus it can be consider as a weakness in our methodology. However, the difference is due to the fact that different extraction methods will also show different sample forms. Therefore, to minimize this case, the researchers weighed all extracts after the evaporation process before preparing the stock solutions. So that the Total Dissolve Solid (TDS) of the extracts in different methods could still be controlled.

CONCLUSION

Sundanese traditional salad (*Lalapan*) contains various bioactive compounds and identified as potentially rich source of antioxidants and antihyperglycemia. The results of phytochemical screening confirms the presence of polyphenols, saponins, tannins, and flavonoids in gandaria leaves, basil, pohpohan, and lettuce. Extraction methods have a significant effect on TPC, TFC, antioxidant activity, and inhibition of α -amylase enzymes. All maceration extracts have higher of TPC, TFC, antioxidants, and inhibition of α -amylase compared to traditional extracts. The leave extract with the highest TPC, TFC, antioxidants, and inhibition of α -amylase activity is gandaria leaves extract with maceration method.

Based on the results of the research, further analysis is needed regarding bioactive compounds in *Lalapan* with different extraction methods using High Performance Liquid Chromatography (HPLC). Pra-clinical and clinical research is needed to determine the benefits of its bioactive

compounds. Furthermore, *Lalapan* as source of antioxidant can be developed as vegetable-based functional food products for antihyperglycemia.

ACKNOWLEDGEMENT

We thank to Ministry of Research, Technology, and Higher Education for providing the research funds so that this study to be part of the main research entitled "Eating Culture, Antioxidant Potential, and Antihyperglycemia of Gandaria Leaves (*Bouea macrophylla* Griff.)" with contracts number 1750/IT3.11/PN/2018 and Dr Joeni Setijo Rahajoe as botanists from Biology Research Center, Cibinong Bogor.

AUTHOR DISCLOSURES

The authors have no conflict of interest.

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